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CLAIMS

What is claimed is:

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- 1. A method of treating glycogen storage disease type II in an individual, deficiency comprising administering to the individual a therapeutically effective amount of human acid α-glucosidase at a regular interval.
- 2. The method of Claim 1, wherein the glycogen storage disease type II is infantile glycogen storage disease type II.
- 3. The method of Claim 1, wherein the glycogen storage disease type II is juvenile glycogen storage disease type II.
- 10 4. The method of Claim 1, wherein the glycogen storage disease type II is adult-onset glycogen storage disease type II.
 - 5. The method of Claim 1, wherein the therapeutically effective amount of human acid α -glucosidase is less than about 15 mg of acid α -glucosidase per kilogram of body weight of the individual.
- 15 6. The method of Claim 5, wherein the therapeutically effective amount of human acid α-glucosidase is about 1-10 mg of acid α-glucosidase per kilogram of body weight of the individual.
 - 7. The method of Claim 5, wherein the therapeutically effective amount of human acid α -glucosidase is about 5 mg of acid α -glucosidase per kilogram of body weight of the individual.

- The method of Claim 1, wherein the human acid α -glucosidase is recombinant haman acid α-glucosidase.
- The method of Claim 1, wherein the human acid α-glucosidase is a precursor of 9. recombinant human acid α -glucosidase.
- 5 The method of Claim-9, wherein the recombinant human acid α -glucosidase is 10. produced in Chinese hamster

- The method of Claim 1, wherein the regular interval is monthly. 11.
- The method of Claim 1, wherein the regular interval is bimonthly. 12.
- The method of Claim 1, wherein the regular interval is weekly. 13.
- The method of Claim 1, wherein the regular interval is twice weekly. 10 14.
 - The method of Claim 1, wherein the regular interval is daily. 15.
 - The method of Claim 1, wherein the human acid α -glucosidase is administered 16. intravenously.
- The method of Claim 1, wherein the human acid α -glucosidase is administered 17. 15 intramuscularly.
 - The method of Claim 1, wherein the human acid α -glucosidase is administered 18. intrathecally or intraventricularly.

- 19. The method of Claim 1, wherein the human acid α-glucosidase is administered in conjunction with an immunosuppressant.
- 20. The method of Claim 19, wherein the immunosuppressant is administered prior to any administration of human acid α-glucosidase to the individual.

Suffs

A method of treating cardiomyopathy associated with glycogen storage disease type II in an individual, comprising administering to the individual a therapeutically effective amount of human acid α -glucosidase at a regular interval.

22. A pharmaceutical composition comprising human acid α-glucosidase in a container with a label containing instructions for administration of the composition for treatment of glycogen storage disease type II.

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